

of HHS to establish a public education program to increase awareness about organ donation and the need to provide for an adequate rate of such donations. In brief, DoT's responsibilities are two-fold: (1) To provide oversight and guidance to the national organ transplant system in the U.S. including monitoring the Organ Procurement and Transplantation Network and the Scientific Registry of Transplant Recipients; and (2) to implement a program of public and professional education and outreach aimed at increasing the number of organ donors in this country. Many preventable deaths occur each year because of a staggering imbalance between the supply and demand for donor organs. As of March 2013, the national transplant waiting list exceeded 117,000. In 2011, the total number of deceased and living organ donors was only 14,145. These donors enabled 28,538 patients to receive a transplant while 6,693 died waiting. Without successful interventions to increase donation, the disparity between need and supply is likely to be substantially

exacerbated, resulting in more unnecessary deaths.

Organdonor.gov is DoT's primary mechanism for providing the public with information about organ donation. Among the most visited pages on organdonor.gov are the donor and recipient life stories which in a recent evaluation study were shown to raise interest on the topic and, more important, persuade people to register as organ donors. To expand this component of organdonor.gov, DoT proposes to develop an application to give organ recipients, living donors, and donor families the opportunity to voluntarily submit their stories to DoT via a standardized online form. The online form will be posted on organdonor.gov and will collect demographic and contact information, the individual's donation/transplant story up to 500 words, a high resolution photo, and a signed authorization. The standardized, electronic form will increase HRSA staff's ability to process those stories more efficiently. In addition to enabling story submission, the online application process will make the donor and recipient life stories posted on the site searchable by the

public to enhance public viewing and understanding of the organ donation process. Submission of a story and completion of the form is voluntary. Overall, this application has the potential to strengthen DoT's outreach efforts and increase organ donation registration in the United States.

*Burden Statement:* Burden in this context means the time expended by persons to generate, maintain, retain, disclose or provide the information requested. This includes the time needed to review instructions; to develop, acquire, install and utilize technology and systems for the purpose of collecting, validating and verifying information, processing and maintaining information, and disclosing and providing information; to train personnel and to be able to respond to a collection of information; to search data sources; to complete and review the collection of information; and to transmit or otherwise disclose the information. The total annual burden hours estimated for this ICR are summarized in the table below.

The annual estimate of burden is as follows:

Form name	Number of respondents	Responses per respondent	Total responses	Hours per response	Total burden hours
Donation/Transplantation Life Story Submission Form .....	100	1	100	0.68	68
Total .....	100	1	100	0.68	68

**ADDRESSES:** Submit your comments to the desk officer for HRSA, either by email to [OIRA\\_submission@omb.eop.gov](mailto:OIRA_submission@omb.eop.gov) or by fax to 202-395-5806. Please direct all correspondence to the "attention of the desk officer for HRSA."

*Deadline:* Comments on this ICR should be received within 30 days of this notice.

Dated: May 7, 2013.

**Bahar Niakan,**  
Director, Division of Policy and Information Coordination.

[FR Doc. 2013-11257 Filed 5-10-13; 8:45 am]

**BILLING CODE 4165-15-P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Health Resources and Services Administration**

**National Advisory Council on Migrant Health; Cancellation of Meeting**

*Name:* National Advisory Council on Migrant Health.

*Dates and Times:* May 21, 2013, 8:30 a.m. to 5:00 p.m., May 22, 2013, 8:00 a.m. to 12:00 p.m.

*Status:* The meeting of the National Advisory Council on Migrant Health, scheduled for May 21 and 22, 2013, is cancelled. This cancellation applies to all sessions of the meeting. The meeting was announced in the **Federal Register** of April 17, 2013 (78 FR 22890).

**FOR FURTHER INFORMATION CONTACT:** Gladys Cate, Public Health Analyst, Office of National Assistance and Special Populations, Bureau of Primary Health Care, Health Resources and Services Administration, 5600 Fishers Lane, Room 15-74, Rockville, Maryland 20857; telephone (301) 594-0367.

Dated: May 7, 2013.

**Bahar Niakan,**  
Director, Division of Policy and Information Coordination.

[FR Doc. 2013-11259 Filed 5-10-13; 8:45 am]

**BILLING CODE 4165-15-P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**National Institutes of Health**

**Proposed Collection; 60-Day Comment Request: National Cancer Institute (NCI) Alliance for Nanotechnology in Cancer Platform Partnership Scientific Progress Reports**

**SUMMARY:** In compliance with the requirement of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, for opportunity for public comment on proposed data collection projects, the National Cancer Institute (NCI), National Institutes of Health (NIH), will publish periodic summaries of proposed projects to be submitted to the Office of Management and Budget (OMB) for review and approval.

Written comments and/or suggestions from the public and affected agencies are invited on one or more of the following points: (1) Whether the proposed collection of information is necessary for the proper performance of the function of the agency, including whether the information will have

practical utility; (2) The accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) Ways to enhance the quality, utility, and clarity of the information to be collected; and (4) Ways to minimize the burden of the collection of information on those who are to respond, including the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology.

*To Submit Comments and for Further Information:* To obtain a copy of the data collection plans and instruments, submit comments in writing, or request more information on the proposed project, contact: Dorothy Farrell, Center for Strategic Scientific Initiatives, Office of Cancer Nanotechnology Research, National Cancer Institute, 31 Center Drive, Bldg. 31 A, Rm. 10A52, Bethesda, MD 20892 or call non-toll-free number 301-496-5652 or Email your request, including your address to: [farrelld@mail.nih.gov](mailto:farrelld@mail.nih.gov). Formal requests for additional plans and instruments must be requested in writing.

*Comment Due Date:* Comments regarding this information collection are best assured of having their full effect if received within 60 days of the date of this publication.

*Proposed Collection:* National Cancer Institute (NCI) Alliance for Nanotechnology in Cancer Platform Partnership Scientific Progress Reports, 0925-NEW, National Cancer Institute (NCI), National Institutes of Health (NIH).

*Need and Use of Information Collection:* National Institutes of Health grantees are required to submit interim and final progress reports and other post-award documents associated with the monitoring, oversight, and closeout of an award. This submission represents a request for OMB to approve new program specific progress report guidelines for Cancer Nanotechnology Platform Partnerships (CNPP) awarded by the National Cancer Institute (NCI). The CNPPs are part of the Alliance for Nanotechnology in Cancer, a network of awards funded by NCI to promote the application of nanotechnology to cancer research and care. The proposed guidelines request information about award performance related to trans-

Alliance collaboration, scientific milestones, progress towards clinical translation and technology commercialization, and education and outreach efforts. The report also gathers information on leveraged funding, patents and publications. The information is gathered every six months. This information is needed to monitor the performance of this special program within NCI, funded through Requests for Applications (RFA CA-09-013, released May 29, 2009) using the cooperative agreement mechanism (U01). The information will be used to monitor individual award performance and the effectiveness of the program as a whole. The respondents are the Principal Investigators of the awards, along with their institutional business officials. The awards are administered by and the reports reviewed by the Office of Cancer Nanotechnology Research (OCNR), part of the Center for Strategic Scientific Initiatives within NCI.

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The estimated annualized burden hours are 72.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Number of respondents	Frequency of response	Average time per response (in hours)	Annual burden hours
Principal Investigators .....	12	2	3	72

Dated: May 7, 2013.  
**Vivian Horovitch-Kelley,**  
*NCI Project Clearance Liaison, NCI, NIH.*  
 [FR Doc. 2013-11294 Filed 5-10-13; 8:45 am]  
**BILLING CODE 4140-01-P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**National Institutes of Health**

**Center for Scientific Review Notice of Closed Meetings**

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. App.), notice is hereby given of the following meetings.

The meetings will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant

applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

*Name of Committee:* Center for Scientific Review Special Emphasis Panel; Member Conflict: Molecular Endocrinology.

*Date:* June 3, 2013.

*Time:* 11:30 a.m. to 1:00 p.m.

*Agenda:* To review and evaluate grant applications.

*Place:* National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892, (Telephone Conference Call).

*Contact Person:* Krish Krishnan, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 6164, MSC 7892, Bethesda, MD 20892, (301) 435-1041, [krishnak@csr.nih.gov](mailto:krishnak@csr.nih.gov).

*Name of Committee:* Vascular and Hematology Integrated Review Group; Hypertension and Microcirculation Study Section.

*Date:* June 6, 2013.

*Time:* 8:00 a.m. to 7:00 p.m.

*Agenda:* To review and evaluate grant applications.

*Place:* Renaissance Mayflower Hotel, 1127 Connecticut Avenue NW., Washington, DC 20036.

*Contact Person:* Ai-Ping Zou, MD, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 4118, MSC 7814, Bethesda, MD 20892, 301-408-9497, [zouai@csr.nih.gov](mailto:zouai@csr.nih.gov).

*Name of Committee:* Immunology Integrated Review Group; Transplantation, Tolerance, and Tumor Immunology Study Section.

*Date:* June 6-7, 2013.

*Time:* 8:00 a.m. to 11:00 a.m.

*Agenda:* To review and evaluate grant applications.

*Place:* Renaissance Washington DC Downtown Hotel, 999 Ninth Street NW., Washington, DC 20001.

*Contact Person:* Jin Huang, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 4199, MSC 7812, Bethesda, MD 20892, 301-435-1230, [jh377p@nih.gov](mailto:jh377p@nih.gov).

*Name of Committee:* Center for Scientific Review Special Emphasis Panel; Health Services Organization and Delivery Overflow.

*Date:* June 11, 2013.

*Time:* 8:00 a.m. to 5:00 p.m.

*Agenda:* To review and evaluate grant applications.